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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 1

Compleat if Known

Application Number	09/853,753
Filing Date	May 14, 2001
First Named Inventor	B ch-Hansen
Group Art Unit	not assigned
Examiner Name	not assigned
Attorney Docket Number	45499.2

OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
OC	AA	Loss of Function Mutations in a Calcium-Channel α_1 subunit gene in Xp11.23 cause incomplete X-linked congenital stationary night blindness - Paper by Bech-Hansen, et al. July, 1998.	
OC	AE	Mutations in NYX, encoding the leucine-rich proteoglycan nyctalopin, cause X-linked incomplete congenital stationary night blindness - Paper by Bech-Hansen et al. November, 2000.	
OC	AC	Evidence for Genetic Heterogeneity in X-linked Congenital Stationary Night Blindness - Paper by Bech-Hansen, et al. - published April 7, 1998.	
OC	AD	Leucine-Rich Repeat Glycoproteins of the Extracellular Matrix - Paper by Hocking, et al. accepted January 29, 1998.	

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